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PATENT

Docket No. 249.00020101

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

Applicant(s):	Dominic E. Cosgrove	Group Art Unit:	1641
Serial No.:	09/970,318	Examiner:	Bao Thuy L. Nguyen
Confirmation No.:	1885		
Filed:	October 3, 2001		

For: IMMUNODIAGNOSTIC DETERMINATION OF USHER SYNDROME TYPE IIA

INFORMATION DISCLOSURE STATEMENT

Assistant Commissioner for Patents
P.O. Box 2327
Arlington, VA 22202

Sir:

In compliance with the duty imposed by 37 C.F.R. § 1.56, and in accordance with C.F.R. §§ 1.97 *et. seq.*, the materials enclosed herewith are brought to the attention of the Examiner as possibly being of interest in connection with the above-identified patent application. Consideration of each of the documents listed on the attached 1449 form(s) is respectfully requested. Pursuant to the provisions of M.P.E.P. §609, Applicant further requests that a copy of the 1449 forms, marked as being considered and initialed by the Examiner, be returned with the next Official Communication.

It is believed that no fee is due, as this Information Disclosure Statement is filed prior to the receipt of any Action on the merits. However, in the event a fee is due, please charge any fee or credit any overpayment to Account No. 13-4895.

The Examiner is invited to contact Applicant's Representatives at the below-listed telephone number, if they can be of any assistance during prosecution of the present application.

CERTIFICATE UNDER 37 C.F.R. 1.8:

The undersigned hereby certifies that this paper is being deposited in the United States Postal Service, as first class mail, in an envelope addressed to: Assistant Commissioner for Patents, P.O. Box 2327, Arlington, VA 22202, on this 13 day of May, 2002.

Ann M. Mueting

May 13, 2002
Date

Respectfully submitted for
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INFORMATION DISCLOSURE STATEMENT	Atty. Docket No.: 249.00020101	Serial No.: 09/970,318
	Applicant(s): Cosgrove	Confirmation No.: 1885
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U.S. PATENT DOCUMENTS

Examiner Initial		Document Number	Date	Name	Class	Subclass	Filing Date If Appropriate
		none					

FOREIGN PATENT DOCUMENTS

Examiner Initial		Document Number	Date	Country	Class	Subclass	Translation	
							Yes	No
		none						

OTHER DOCUMENTS (Including Authors, Title, Date, Pertinent Papers, etc.)

Examiner Initial		Document Description
		Adato et al., "Three novel mutations and twelve polymorphisms identified in the USH2A gene in Israeli USH2 families," *Hum Mutat. (Mutation in Brief)*; published online for subscribers 2000 Mar 22 (6 pgs.).
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		Barkalow et al., "Localization of the major heparin-binding site in fibronectin," *J Biol Chem.* 1991 Apr 25;*266*(12):7812-8.
		Beck et al., "Structure and function of laminin: anatomy of a multidomain glycoprotein," *FASEB J.* 1990 Feb 1;*4*(2):148-60.
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		Boughman et al., "Usher syndrome: definition and estimate of prevalence from two high-risk populations," *J Chronic Dis.* 1983;*36*(8):595-603.
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EXAMINER	Date Considered

*Examiner: Initial if citation considered, whether or not citation is in conformance with MPEP 609; Draw line through citation if not in conformance and not considered. Include copy of this form with next communication to applicant.

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Examiner Initial		Document Description
		Brookhouser, Patrick E., "Center for Hearing Loss in Children," Grant Abstract, Grant Number 5P60DC000982-10 [online]. National Institutes of Health, National Institute on Deafness and other Disorders of Communication, project dates 1990-09-30 to 2001-08-31. Retrieved from the Internet 2002-03-21. URL: <http://commons.cit.nih.gov/crisp3/CRISP_LIB.getdoc?textkey=6055813&p_grant_num=5P60DC000982-10&p_query=&ticket=388948&p_audit_session_id=3084290&p_keywords=>; 2 pages.
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		Kimberling et al., "Gene mapping of Usher syndrome type IIa: localization of the gene to a 2.1-cM segment on chromosome 1q41," *Am J Hum Genet.* 1995 Jan; *56*(1):216-23.
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		National Center for Biotechnology Information, National Library of Medicine, National Institutes of Health, GenBank Locus HSUSH2A01, Accession No. AF091873, "Homo sapiens Usher syndrome type IIa protein gene, exons 1 and 2," [online]. Bethesda, MD [retrieved on 2002-04-05]. Retrieved from the Internet: <URL:http://www.ncbi.nlm.nih.gov/entrez/viewer.fcgi?query_key=4&db=nucleotide&page=0&dispmax=20&WebEnv=Wg%3Cn_FDG%5DE%60%3E%3D%3Cc%5DPGDJc_gTB%5EjbFkl%3C_JEH%3Dzcc%3EF%5EfFJdTI%3D%3F%3C&WebEnvRq=1>; 3 pgs.

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Examiner Initial		Document Description
		National Center for Biotechnology Information, National Library of Medicine, National Institutes of Health, GenBank Locus HSUSH2A17, Accession No. AF091889, "Homo sapiens Usher syndrome type IIa protein gene, exon 21 and complete cds," [online]. Bethesda, MD [retrieved on 2002-04-05]. Retrieved from the Internet: <URL: http://www.ncbi.nlm.nih.gov/entrez/viewer.fcgi?query_key=6&db=nucleotide&page=0&dispmax=20&WebEnv=C%5B%60%3CIgCAA%5C%5Ee%3D%3FHCK%3CCDB%3CdlEjhCdbaiH%3F%40AARqjH%5EGe%5E%3EkDFH&WebEnvRq=1>; 3 pgs.
		Nuutila, "Dystrophia retinae pigmentosa–dysacusis syndrome (DRD): a study of the Usher- or Hallgren syndrome," *J Génét Hum.* 1970 May; *18*(1):57-88.
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		van der Loop et al., "Identification of COL4A5 defects in Alport's syndrome by immunohistochemistry of skin," *Kidney Int.* 1999 Apr;*55*(4):1217-24.
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		Weston et al., "Genomic structure and identification of novel mutations in usherin, the gene responsible for Usher syndrome type IIa," *Am J Hum Genet.* 2000 Apr;*66*(4):1199-210.
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DOCUMENTS ACCOMPANYING INFORMATION DISCLOSURE STATEMENT FILED MAY 13, 2002

PART 1 OF 2

PATENT
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PART 2 OF 2